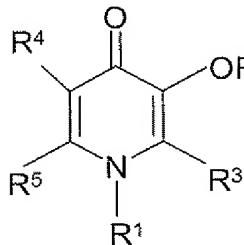
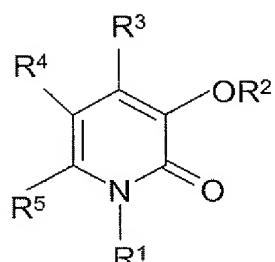


Claims:

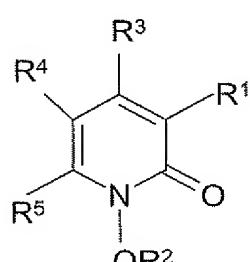
1. **(Currently amended)** A method for treating a skin microcirculatory disorder (SMD) comprising topically administering a hydroxypyridonone of formulae (I-III):



(I)



(II)



(III)

wherein

R<sup>1</sup> represents a (C<sub>1</sub>-C<sub>10</sub>)- alkyl, (C<sub>1</sub>-C<sub>10</sub>)-alkenyl, (C<sub>1</sub>-C<sub>10</sub>)-alkoxy, (C<sub>1</sub>-C<sub>10</sub>) hydroxyalkyl, (C<sub>5</sub>-C<sub>12</sub>)-aralkyl, (C<sub>3</sub>-C<sub>12</sub>)-cycloalkyl, (C<sub>1</sub>-C<sub>8</sub>)- carboalkoxy or (C<sub>1</sub>-C<sub>8</sub>)- carbamyl, or a (C<sub>10</sub>-C<sub>30</sub>)-peptide or peptidomimetic moiety, or a (C<sub>3</sub>-C<sub>6</sub>) polyol or monosaccharide;

R<sup>2</sup> represents an hydrogen atom or a linear or branched, saturated or unsaturated to (C<sub>1</sub>-C<sub>22</sub>)-acyl, optionally substituted by (C<sub>1</sub>-C<sub>8</sub>)-alkoxy, carboxy, (C<sub>1</sub>-C<sub>8</sub>) alkoxy carbonyl, amino, hydroxy, said amino and hydroxy being optionally (C<sub>1</sub>-C<sub>22</sub>)-acylated or - alkylated;

R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup>, each individually, represent a hydrogen atom, or (C<sub>1</sub>-C<sub>10</sub>)-alkyl, (C<sub>1</sub>-C<sub>10</sub>)- alkenyl, (C<sub>1</sub>-C<sub>10</sub>)-alkoxy, (C<sub>5</sub>-C<sub>12</sub> aryl) alkyl, (C<sub>5</sub>-C<sub>12</sub>)-cycloalkyl, (C<sub>1</sub>-C<sub>8</sub> carbo)-alkoxy or (C<sub>1</sub>-C<sub>8</sub>)- carbamyl group;

with the proviso that both R<sup>1</sup> and R<sup>3</sup> are not hydrogen;

or a dermatologically/cosmetically salt thereof.

2. **(Previously presented)** A method according to claim 1, wherein the skin microcirculatory disorder (SMD) is rosacea.

3. **(Previously presented)** A method according to claim 1, wherein the skin microcirculatory disorder (SMD) is cutaneous vasculitis.

4. **(Previously presented)** A method according to claim 1, wherein the skin microcirculatory disorder (SMD) is actinic purpura.

5. (Previously presented) A method according to claim 1, wherein the skin microcirculatory disorder (SMD) is a skin capillaritis.

**6.(Currently amended)** A method according to claim 8, wherein the skin capillaritis is progressive pigmentary dermatosis, purpura annularis telangiectodes, ~~lichen aureus~~, contact allergy skin capillaritis, lichens aureus, itching purpura, or eczematid-like purpura, or pigmented purpuric lichenoid dermatosis.

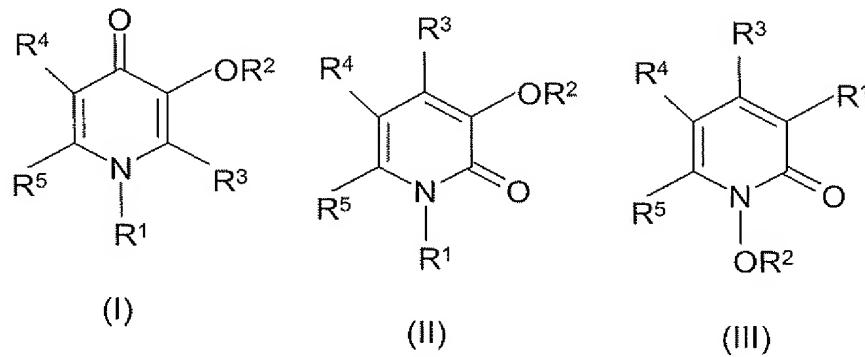
7. (Cancelled)

8. (Withdrawn) A method according to claim 1, wherein R<sup>1</sup> and R<sup>2</sup> are methyl, R<sup>3</sup> and R<sup>4</sup> are hydrogens.

9. (Withdrawn) A method according to claim 1, wherein R<sup>1</sup> and R<sup>2</sup> are ethyl R<sup>3</sup> and R<sup>4</sup> are hydrogens.

10. (Withdrawn) A method according to claim 1, wherein R<sup>1</sup> is CH<sub>2</sub>CH<sub>2</sub>OH, R<sup>2</sup> is methyl or ethyl, and R<sup>3</sup> and R<sup>4</sup> are hydrogens.

11. **(Currently amended)** A method for the treatment of skin microcirculatory disorder (SMD) comprising locally applying to a mammal in need thereof of a therapeutically effective amount of hydroxypyridonone compound of formulae (I-III):



wherein

$R^1$  represents a (C<sub>1</sub>-C<sub>10</sub>)- alkyl, (C<sub>1</sub>-C<sub>10</sub>)-alkenyl, (C<sub>1</sub>-C<sub>10</sub>)-alkoxy, (C<sub>1</sub>-C<sub>10</sub>) hydroxyalkyl, (C<sub>5</sub>-C<sub>12</sub>)- aralkyl, (C<sub>3</sub>-C<sub>12</sub>)-cycloalkyl, (C<sub>1</sub>-C<sub>8</sub>)- carboalkoxy or (C<sub>1</sub>-C<sub>8</sub>)- carbamyl, or a (C<sub>10</sub>-C<sub>30</sub>)- peptide or peptidomimetic moiety, or a (C<sub>3</sub>-C<sub>6</sub>) polyol or monosaccharide;

$R^2$  represents an hydrogen atom or a linear or branched, saturated or unsaturated ( $C_1$ - $C_{22}$ )-acyl, optionally substituted by ( $C_1$ - $C_8$ )-alkoxy, carboxy, ( $C_1$ - $C_8$ ) alkoxy carbonyl, amino, hydroxy, said amino and hydroxy being optionally ( $C_1$ - $C_{22}$ )-acylated or - alkylated;

$R^3$ ,  $R^4$  and  $R^5$ , each individually, represent a hydrogen atom, or  $(C_1-C_{10})$ -alkyl,  $(C_1-C_{10})$ - alkenyl,  $(C_1-C_{10})$ -alkoxy,  $(C_5-C_{12})$  aryl alkyl,  $(C_5-C_{12})$ -cycloalkyl,  $(C_1-C_8)$  carbo)-alkoxy or  $(C_1-C_8)$ - carbamyl group;

with the proviso that both  $R^1$  and  $R^3$  are not hydrogen;

or a dermatologically/cosmetically acceptable salt thereof

in admixture with a dermatologically/cosmetically acceptable carrier.

12. **(Previously presented)** A method according to claim 11, for the treatment of rosacea, cutaneous vasculitis, or actinic purpura.

13. **(Currently amended)** A method according to Claim 14 11, for the treatment of progressive pigmented purpura, itching purpura, pigmented purpuric lichenoid dermatosis, purpura annularis telangiectodes, or contact allergy skin capillaritis, or lichens aureus.

14. **(Currently amended)** A method according to Claim 14 11, for the treatment of traumatic skin haemorrhage, a complication of sclerotherapy, lipoplasty or tattooing, drug induced pigmented purpuric dermatosis, or actinic purpura.

15. **(Withdrawn)** A method according to claim 11, wherein  $R^2$ ,  $R^3$ ,  $R^4$  and  $R^5$ , each individually, represent a hydrogen atom.

16. **(Previously presented)** A method according to claim 11, wherein  $R^1$  and  $R^3$  each individually, represent  $(C_1-C_4)$ - alkyl, hydroxyalkyl or alkoxy.

17. **(Withdrawn)** A method according to claim 11, wherein said  $R^2$  acyl group is formed by unbranched, naturally occurring caprylic acid, cupric acid, lauric acid, myristic acid, palmitic acid, palmitoleic acid, stearic acid, oleic acid, vaccenic, linoleic acid, alpha-linolenic acid, eleostearic, delta-linolenic acid, gondoic acid, dihomo- $\gamma$ -linolenic acid, arachidonic acid, eicosapentaenoic acid, docosenoic acid, docosatekaenoic acid, docosapentaenoic acid, docosapentaenoic, docosahexacuoic acid, nervonic or a mixture thereof.

18. **(Withdrawn)** A method according to claim 11, wherein said  $R^2$  acyl is a  $C_{1-8}$  which is branched at the carbon atom adjacent to the carbonyl group.

19. **(Currently amended)** A method according to claim 11, wherein said hydroxypyridonone is 1, 2 dimethyl-3-hydroxy-4-pyridinone (deferiprone); 1,2-diethyl-3- hydroxy- 4-pyridinone; 1-methyl-2-ethyl-3-hydroxy-4-pyridinone; ~~1-methyl-2-ethyl-3-hydroxy-4-pyridinone~~ or 1-methyl-2-(2-methoxy-ethyl)-3-hydroxy-4- pyridinone.